CHRNG gene

cholinergic receptor nicotinic gamma subunit

Normal Function

The *CHRNG* gene provides instructions for making the gamma (γ) protein component (subunit) of the acetylcholine receptor (AChR) protein. The AChR protein is found in the membrane of skeletal muscle cells and is critical for signaling between nerve and muscle cells. Signaling between these cells is necessary for movement. The AChR protein consists of five subunits, each of which is produced from a different gene. The subunits are assembled into the AChR protein in the endoplasmic reticulum, a cell structure involved in protein processing and transport, before being transported to the cell membrane. The γ subunit is found only in the fetal AChR protein. At about the thirty-third week of pregnancy, the γ subunit is replaced by the epsilon (ϵ) subunit, which is produced by the *CHRNE* gene, to form the adult AChR protein.

Health Conditions Related to Genetic Changes

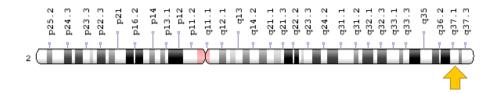
multiple pterygium syndrome

At least 14 mutations in the *CHRNG* gene have been found to cause multiple pterygium syndrome, a condition characterized by webbing of the skin (pterygium) and a lack of muscle movement (akinesia) before birth. These mutations include replacing, adding, or deleting DNA building blocks (nucleotides). *CHRNG* gene mutations result in an impaired or missing γ subunit. The severity of the *CHRNG* gene mutation influences the severity of the condition. Typically, mutations that prevent the production of any γ subunit will result in lethal multiple pterygium syndrome, which is fatal before birth, while mutations that allow the production of some γ subunit will lead to the milder form of this condition called multiple pterygium syndrome, Escobar type. A shortage of a functional γ subunit prevents the fetal AChR protein from being assembled or properly placed in the muscle cell membrane. As a result, the fetal AChR protein cannot function and the communication between nerve and muscle cells in the developing fetus is impaired. A lack of signaling between nerve and muscle cells leads to akinesia and pterygium before birth, and many of the other signs and symptoms of multiple pterygium syndrome.

Chromosomal Location

Cytogenetic Location: 2q37.1, which is the long (q) arm of chromosome 2 at position 37.1

Molecular Location: base pairs 232,539,727 to 232,546,328 on chromosome 2 (Homo sapiens Annotation Release 108, GRCh38.p7) (NCBI)



Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- acetylcholine gamma muscle receptor subunit
- acetylcholine receptor subunit gamma
- acetylcholine receptor, muscle, gamma subunit
- ACHG_HUMAN
- ACHRG
- cholinergic gamma nicotinic receptor
- cholinergic receptor, nicotinic gamma
- cholinergic receptor, nicotinic, gamma
- cholinergic receptor, nicotinic, gamma (muscle)
- cholinergic receptor, nicotinic, gamma polypeptide

Additional Information & Resources

Educational Resources

- Molecular Biology of the Cell (fourth edition, 2002): The Acetylcholine Receptors at the Neuromuscular Junction Are Transmitter-gated Cation Channels https://www.ncbi.nlm.nih.gov/books/NBK26910/#A2055
- Molecular Cell Biology (fourth edition, 2000): All Five Subunits in the Nicotinic Acetylcholine Receptor Contribute to the Ion Channel https://www.ncbi.nlm.nih.gov/books/NBK21586/#A6233
- Molecular Cell Biology (fourth edition, 2000): Signals From the Motor Neuron Regulate the Expression and Localization of the Acetylcholine Receptor (AChR) (figure) https://www.ncbi.nlm.nih.gov/books/NBK21742/figure/A6882/

Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28CHRNG%5BTIAB%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D

OMIM

 CHOLINERGIC RECEPTOR, NICOTINIC, GAMMA POLYPEPTIDE http://omim.org/entry/100730

Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology http://atlasgeneticsoncology.org/Genes/GC_CHRNG.html
- ClinVar https://www.ncbi.nlm.nih.gov/clinvar?term=CHRNG%5Bgene%5D
- HGNC Gene Family: Cholinergic receptors nicotinic subunits http://www.genenames.org/cgi-bin/genefamilies/set/173
- HGNC Gene Symbol Report http://www.genenames.org/cgi-bin/gene_symbol_report?q=data/ hgnc_data.php&hgnc_id=1967
- NCBI Gene https://www.ncbi.nlm.nih.gov/gene/1146
- UniProt http://www.uniprot.org/uniprot/P07510

Sources for This Summary

- OMIM: CHOLINERGIC RECEPTOR, NICOTINIC, GAMMA POLYPEPTIDE http://omim.org/entry/100730
- Hoffmann K, Muller JS, Stricker S, Megarbane A, Rajab A, Lindner TH, Cohen M, Chouery E, Adaimy L, Ghanem I, Delague V, Boltshauser E, Talim B, Horvath R, Robinson PN, Lochmüller H, Hübner C, Mundlos S. Escobar syndrome is a prenatal myasthenia caused by disruption of the acetylcholine receptor fetal gamma subunit. Am J Hum Genet. 2006 Aug;79(2):303-12. Epub 2006 Jun 20.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16826520
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1559482/
- Michalk A, Stricker S, Becker J, Rupps R, Pantzar T, Miertus J, Botta G, Naretto VG, Janetzki C, Yaqoob N, Ott CE, Seelow D, Wieczorek D, Fiebig B, Wirth B, Hoopmann M, Walther M, Körber F, Blankenburg M, Mundlos S, Heller R, Hoffmann K. Acetylcholine receptor pathway mutations explain various fetal akinesia deformation sequence disorders. Am J Hum Genet. 2008 Feb;82(2): 464-76. doi: 10.1016/j.ajhg.2007.11.006.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18252226
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2427255/
- Morgan NV, Brueton LA, Cox P, Greally MT, Tolmie J, Pasha S, Aligianis IA, van Bokhoven H, Marton T, Al-Gazali L, Morton JE, Oley C, Johnson CA, Trembath RC, Brunner HG, Maher ER. Mutations in the embryonal subunit of the acetylcholine receptor (CHRNG) cause lethal and Escobar variants of multiple pterygium syndrome. Am J Hum Genet. 2006 Aug;79(2):390-5. Epub 2006 Jun 20.

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